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(21) International Application Number: PCT/IT96/00097			(74) Agents: DE SIMONE, Domenico et al.: Ing. Barzano' & Zanardo Roma S.p.A., Via Piemonte, 26, I-00187 Roma (IT).
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(71) Applicants (for all designated States except US): UNIVERSITA' DEGLI STUDI DI ROMA "LA SAPIENZA" [IT/IT]; Piazzale Aldo Moro, 5, I-00185 Roma (IT). ISTITUTO SUPERIORE DI SANITA [IT/IT]; Viale Regina Elena, 299, I-00161 Roma (IT).			
(72) Inventors; and (75) Inventors/Applicants (for US only): CASSONE, Antonio [IT/IT]; Istituto Superiore di Sanita', Viale Regina Elena, 299, I-00161 Roma (IT). LA VALLE, Roberto [IT/IT]; Istituto Superiore di Sanita', Viale Regina Elena, 299, I-00161 Roma (IT). BROMURO, Carla [IT/IT]; Istituto Superiore di Sanita', Viale Regina Elena, 299, I-00161 Roma (IT). CRISANTI, Andrea [IT/IT]; Universita' degli Studi di Roma "La Sapienza", Piazzale Aldo Moro, 5, I-00185 Roma (IT). MULLER, Hans, Michael [IT/IT]; Universita' degli Studi di Roma "La Sapienza", Piazzale Aldo Moro, 5, I-00185 Roma (IT).			
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Published*With international search report.***(54) Title:** CANDIDA HEAT SHOCK PROTEIN, cDNA AND USES THEREOF**(57) Abstract**

A nucleotide sequence and related protein from Candida homologous to 70 kd heat shock protein, for uses in diagnosis and therapy.

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INTERNATIONAL SEARCH REPORT

International application No.

/IT 96/ 00097

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:
Please see Futher Information sheet enclosed.
2. ☐ Claims Nos.:
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.

CANDIDA HEAT SHOCK PROTEIN, CDNA AND USES THEREOF

The invention concerns the cDNA and the corresponding protein of a heat shock protein isolated from C. albicans, and fragments thereof to develop
5 methods to identify C. albicans in biological and/or environment samples, and/or preparations either for therapeutic, prophylaxis or vaccine purpose.

Pathogenic yeasts are the major agents of opportunistic infections in immunosuppressed patients, in
10 particular AIDS, tumor, neutropenia patients or bone marrow transplanted subjects (1). HIV⁺ subject susceptibility to C. albicans is related to the strong decrease of cell-mediated immunity because of the numerical and functional decrease of CD4⁺ helper-inducer
15 lymphocytes (2).

C. albicans cell wall mannoproteins and heatshock proteins of other microorganisms as well, are major antigens and immunomodulators, and play a relevant role during host invasion and infection (3,4).

20 By using a rabbit immune serum obtained against heat-inactivated C. albicans ATCC 20955 strain cells, the authors of the instant invention isolated the caRLV130 clone from an expression library in the λ gt11 phage obtained by cDNA isolated from C. albicans at the yeast
25 growth stage. Said clone contains a DNA insert of 2325 base pairs which codes in the 5'-3' direction from +105 to +2072 for a 656 aminoacid protein having a strong homology with a S. cerevisiae heat shock protein 70.

30 HSPs are induced by different stresses, either chemical or physical, normally by heating. Many HSPs are present and active also in non stressed cells, where they play important functions of cell physiology ("chaperonins"). They may be grouped in families of

different molecular weights, very conserved even among phylogenesis distant organisms (5). Therefore it should not be surprising either that HSPs are involved in the immune response, or that they represent major antigens of different pathogenic agents, or that they may give autoimmune responses, given to the fact that the infection itself represents an extreme form of stress, both for the infectious agent and for the host (4).

It is therefore an object of the invention a nucleic acid comprising a nucleotide sequence coding the protein having the amino acid sequence of SEQ ID No.2 or parts thereof. Preferably the nucleic acid comprises a nucleotide sequence with at least a 65% homology with the nucleotide sequence of SEQ ID No.1 or parts thereof. More preferably the nucleic acid comprises the nucleotide sequence of SEQ ID No.1 or parts thereof.

Further object of the invention is a composition comprising a nucleic acid comprising a nucleotide sequence coding the protein having the amino acid sequence of SEQ ID No.1 or parts thereof. Preferably the composition comprises a nucleic acid having a nucleotide sequence with at least a 65% homology with the nucleotide sequence of SEQ ID No.1 or parts thereof. More preferably the composition comprises a nucleic acid having the nucleotide sequence of SEQ ID No.1 or parts thereof.

Further object of the invention is the use of the nucleic acid comprising a nucleotide sequence coding the protein having the amino acid sequence of SEQ ID No.2 or parts thereof for oligonucleotide probes to be used in diagnosis and typing of Candida related pathologies. The use of a nucleic acid having at least a 65% homology with the nucleotide sequence of SEQ ID No.1 or parts thereof is preferred. The use of a nucleic acid having the

nucleotide sequence of SEQ ID No.1 or parts thereof is most preferred.

The oligonucleotides of the invention are advantageously used for PCR (polymerase chain reaction) to detect the presence in biological and/or environment samples either of C. albicans or of other Candida species or of yeast-like related microorganisms comprising said gene; in a labeled form (radionuclides, biotin, enzymes, etc.) to detect the presence in biological and/or environment samples either of C. albicans or of other related; for the C. albicans or related species typing and/or diagnosis; as potential antibiotic and/or chemiotherapeutic targets, or antisense RNA active for Candida species and/or yeast-like related microorganisms coding an homologous sequence.

Another object of the invention is a polypeptide having the aminoacid sequence comprised in the SEQ ID No.2, or having at least a 50% homology with SEQ ID No. 2 or fragments, and/or functional and immunologic homologous thereof.

Further object of the invention is a composition comprising a polypeptide having an amino acid sequence comprised in SEQ ID No.2 or having at least a 50% homology with SEQ ID No. 2 or fragments, and/or functional and immunologic homologous thereof.

Further object of the invention is the use of a polypeptide having the amino acid sequence comprised in SEQ ID No.2 or having at least a 50% homology with SEQ ID No. 2 or of fragments, and/or functional and/or immunologic homologous thereof to make polyclonal or monoclonal antibodies against the 70 kd heat shock protein (HSP70) of C. albicans or related species.

Further object of the invention is the use of a polypeptide having the amino acid sequence comprised in

SEQ ID No.2 or having at least a 50% homology with SEQ ID No. 2 or of fragments, and/or functional and/or immunologic homologous thereof to detect C. albicans and related species HSP70 in a biological sample having a human, animal or environmental origin.

Further object of the invention is the use of a polypeptide having the amino acid sequence comprised in SEQ ID No.2 or having at least a 50% homology with SEQ ID No. 2 or of fragments, and/or functional and/or immunologic homologous thereof for the preparation of a composition to be used for prophylaxis and/or therapy of C. albicans or related microorganisms (pathogenic yeasts) diseases.

Further object of the invention is the use of a polypeptide having the amino acid sequence comprised in SEQ ID No.2 or having at least a 50% homology with SEQ ID No. 2 or of fragments, and/or as potential antibiotic and/or chemiotherapeutic targets active for Candida species and/or yeast-like related microorganisms coding an homologous sequence.

The invention will be described in different embodiments for clarifying but not limiting purposes.

Figure 1 represents the 1971 base pair DNA sequence (small letters) corresponding to the open reading frame of λ gt11-(caRLA130) clone insert and deduced aminoacid sequence (capital letters one-letter code).

Figure 2 represents the nucleotide sequence of the coding insert of caRLV130 clone (small letter) and comparison with S. cerevisiae YSCSSA1 gene (capital letter).

Figure 3 represents the 656 aminoacid sequence deduced from the coding insert of caRLV130 clone (small letter) and comparison with the S. cerevisiae YSCSSA1

gene (capital letter). The aminoacid code utilized is the one letter code.

Figure 4 represents in panel A. Southern blot analysis of C. albicans strain ATCC 20955 chromosomes, obtained by pulse field electrophoresis (TAFE). The caRLV130 probe labeling refers to the highest molecular weight chromosome (3.5 Mbp). In panel B. Electrophoretic separation of C. albicans strain ATCC 20955 chromosomes.

Figure 5 represents on the left side: Northern blot analysis by hybridization of total RNA extracted from C. albicans cells grown at 22°C and transferred at 37°C for the time indicated with radiolabeled caRLV130 (cahsp70) and actin probes. The actin probe hybridization was performed to control the RNA amount on filters (see ref. 8). On the right side: immunoblotting reactivity of anti-CAHSP70 mouse serum with C. albicans extracts, at different times further to inducing a heat shock response as previously described.

Figure 6 represents in panel A. SDS-PAGE analysis: a) expression products of E. coli M15 containing the pDS56/RBS-E⁻-6his caRLV130/1 plasmid; b) expression products of E. coli M15 containing the pDS56/RBS-E⁻-6his caRLV130/2 plasmid; c) expression products of E. coli M15 containing the pDS56/RBS-E⁻-6his caRLV130/3 plasmid; d) expression products of E. coli M15 containing the pDS56/RBS-E⁻-6his caRLV130/4 plasmid. N.I.: Non induced E. coli culture extracts. I.: 1 mM IPTG induced E. coli culture extracts. P.: Purified fraction on histidine affinity nickel column from 1 mM IPTG induced E. coli culture extracts. In panel B. Schematic representation of caRLV130 coding sequence portions cloned into recombinant plasmids used in panel A. Right side: molecular weight in kDa. Left side: denomination of the expression product of recombinant plasmid. For further details, see table I.

Figure 7 represents the reactivity after immunoblotting on nitrocellulose filters of mouse sera as shown in the figure obtained against CAHSP70 fragments; a) expression products of nickel column purified pDS56/RBSII-E⁻-6his caRLV130/1 plasmid in 1 mM IPTG induced E. coli; b) expression products of nickel column purified pDS56/RBSII-E⁻-6his caRLV130/2 plasmid in 1 mM IPTG induced M15 E. coli; c) expression products of nickel column purified pDS56/RBSII-E⁻-6his caRLV130/3 plasmid in 1 mM IPTG induced M15 E. coli; d) expression products of nickel column purified pDS56/RBSII-E⁻-6his caRLV130/4 plasmid in 1 mM IPTG induced M15 E. coli (see also Fig. 6 and table I for a definition of polypeptide fragments). Left side: molecular weight of purified fragments.

Figure 8 represents the reactivity after immunoblotting on nitrocellulose filters of wealthy human sera obtained against CAHSP70 and fragments thereof; a) expression products of nickel column purified pDS56/RBSII-E⁻-6his caRLV130/1 plasmid in 1 mM IPTG induced M15 E. coli; b) expression products of nickel column purified pDS56/RBSII-E⁻-6his caRLV130/2 plasmid in 1 mM IPTG induced M15 E. coli; c) expression products of nickel column purified pDS56/RBSII-E⁻-6his caRLV130/3 plasmid in 1 mM IPTG induced M15 E. coli; d) expression products of nickel column purified pDS56/RBSII-E⁻-6his caRLV130/4 plasmid in 1 mM IPTG induced M15 E. coli. Left side: molecular weight of purified fragments. Right side: denomination of purified protein fragments. For further details see also table I.

Figure 9 represents in panel A. PCR experiment performed using oligonucleotide combination CA2-CA3 in the presence of C. albicans, C. parapsilosis (2), C. glabrata (3), C. guilliermondii (4), C. krusei (5), C.

tropicalis (6), Mus muris (7), E. coli (8), S. cerevisiae (9) DNAs. Control with no DNA is as (10). At the right side the molecular weight of the amplified fragment is indicated. In panel B. PCR experiment using the combination of CA1-CA4 oligonucleotides in the presence of C. albicans cDNA: DNA amplified from C. albicans DNA: 10 ng (2); 1 ng (3); 100 pg (4); 10 pg (5); 1 pg (6). Control: reaction with no DNA (1). PCR reaction conditions are as follows: 90 sec. 94°C denaturation; 90 sec. 60°C annealing; 120 sec. 72°C extension; 25 cycles.

Figure 1 shows the 1971 bp coding region of the isolated gene.

The caRLV130 sequence was filed with EMBL data base (No. Z30210). No intron can be found in the intronic sequence, as shown by PCR product analysis and by "Southern-blot". By comparing the caRLV130 insert sequence with sequences present in the 6.7 version "GENE BANK" data base, some homologies can be detected. The insert shows the most high homology with the S. cerevisiae gene SSA1 (one of the nine heatshock yeast gene family). The overall nucleotide sequence homology is of 78.8% in the coding region (figg. 2 and 3).

The gene corresponding to the caRLV130 sequence was mapped on the C. albicans chromosome showing the highest molecular weight (3.5 Mpb) by pulse field electrophoresis (transverse-alternate: TAFE) utilizing the caRLV130 labeled cDNA insert as hybridization probe with C. albicans chromosomes blotted on nitrocellulose filters (fig. 4A and 4B). Gene transcription is activated by exposing cells to a temperature higher than room temperature (thermal shift from 22°C to 37°C). Such finding was demonstrated by hybridization experiments using C. albicans total RNA (from cells grown either at 22°C or at 37°C, fractionated according to molecular

weight on formaldehyde agarose gel and blotted on nitrocellulose filters) and the caRLV130 DNA insert as radioactive probe. The induction of transcription is coupled also to an increase of protein expression, 2, 6
5 and 24 hours further to the 22°C to 37°C temperature shift (see fig. 5).

Different portions of the caRLV130 insert sequence were cloned in the expression plasmid pDS56/RBSII-E⁻-6his (6), and coded polypeptides were expressed in E. coli
10 after fusion of their amino terminal sequence with 6 histidine residues. The histidine stretch allowed to a rapid and efficient purification of polypeptides derived from the caRLV130 insert sequence on nickel columns (see
15 fig. 6 and table I for denomination and length of polypeptide fragments).

Table I

Definition of nickel column purified CAHSP70 polypeptides. Peptides are coded by pDS56/RBSII-E⁻6his recombinant plasmids wherein caRLV130 fragments were cloned. The position refers to nucleotide and aminoacid sequences as shown in Fig. 1. The peptide length refers to the fusion product coded by the recombinant plasmid.

coding DNA				coded peptide			
denomination	sequence location on cDNA	position (nt)	length (bp)	denomination	peptide fragment location	position (aa)	length
caRLV130/1	whole coding	1-2229	2229	CAHSP70	whole protein	1-656	71.3 664
caRLV130/2	3' end cDNA	1393-2229	837	CAHSP70/2	C-terminus	465-656	21.0 202
caRLV130/3	5' end cDNA	1-732	732	CAHSP70/3	N-terminus	1-244	28.4 261
caRLV130/4	5' end cDNA	1-1027	1027	CAHSP70/4	N-terminus	1-342	39.4 358

After purification, recombinant peptides were used as immunogens to produce mouse immune sera and are therefore able also to induce monoclonal antibodies. Therefore, according the immunization schedule shown in table II, polypeptides, and the whole purified protein as well, induce specific antibodies in a 18-22 g weight Balb/c mouse.

Table II

Immunization schedule of 18-22 g weight Balb/c mice with CAHSP70 peptides purified as described in the text and in Fig.6.

Immunogen	Immunization (day 1)	First boost (day 21)	Second boost (day 41)	Serum titer (day 51)
CAHSP70	5 µg	5 µg	10 µg	> 12.800
CAHSP70/2	5 µg	5 µg	10 µg	> 12.800
CAHSP70/3	5 µg	5 µg	10 µg	> 12.800
CAHSP70/4	5 µg	5 µg	10 µg	> 12.800

The indicated immunogen concentration was inoculated intraperitoneally in a 200 µl volume. The titer was determined by indirect ELISA with the antigen used for coating at a 200 ng/well concentration, in a final volume of 100 µl, and represents the highest serum dilution able to give an ELISA positive reaction (optical density at 405 nm \geq two fold the no antigen control value).

Serum titers for each antigen resulted to be > 12.800 by immunoenzyme test (indirect ELISA) with the adsorbed antigen at 200 ng/well, in a final volume of 100 µl. The specificity of immunoenzyme test results were confirmed in immunoblot experiments on nitrocellulose filters, as shown in Fig. 7.

The same polypeptides were utilized as immunogens in proliferation assays on peripheral human blood lymphocytes by evaluating the ^3H -thymidine uptake further to 7 day culturing according to standard techniques (7).

Results obtained with different donors (two examples are shown in table III) demonstrate that CAHSP70 is able to induce a good thymidine uptake and the proliferation of naive lymphocytes from umbilical cord blood (Table IV), suggesting that the protein itself or parts thereof has a mitogenic activity.

Table III

Peripheral blood lymphocytes proliferation induction activity of CAHSP70 and fragments thereof

10

inducing materials	dose	lymphoproliferative activity ³ H-thymidine uptake (cpm \pm SD/2x10 ⁴ cells)
none	-	500 \pm 200
MP-F2	50 μ g/ml	13.393 \pm 11.555
IL-2	100 U/ml	28.205 \pm 18.014
CAHSP70	1 μ g/ml	8.730 \pm 5.181
CAHSP70/2	1 μ g/ml	2.900 \pm 2.300
CAHSP70/3	1 μ g/ml	3.600 \pm 2.700
CAHSP70/4	1 μ g/ml	11.685 \pm 8.174

Lymphoproliferation of wealthy donor peripheral blood lymphomonocyte cultures further to induction with the CAHSP70 cloned fragments. Positive controls: C. albicans mannoproteic antigen (MP-F2) and Interleukin-2 (IL-2). Negative controls: no materials. Shown values represent average values \pm SD from 7 experiments with 5 different donors. ³H-thymidine uptake was determined after 7 days of culture. For technical details, see ref. 7.

15

Table IV

Umbilical cord blood cell proliferation induction
activity of CAHSP70 and fragments thereof

inducing materials	dose	lymphoproliferative activity ³ H-thymidine uptake (cpm \pm SD/2x10 ⁵ cells)	
		cord blood 1	cord blood 2
none		2.5 \pm 0.4	1.3 \pm 0.3
IL-2	100 U/ml	37.7 \pm 4.5	32.8 \pm 6.0
MP-F2	50 μ g/ml	3.0 \pm 1.4	1.5 \pm 0.4
CAHSP70	1 μ g/ml	12.5 \pm 1.8	22.8 \pm 6.6
CAHSP70/2	1 μ g/ml	18.2 \pm 3.0	23.1 \pm 3.9
CAHSP70/3	1 μ g/ml	23.8 \pm 5.4	20.6 \pm 9.2
CAHSP70/4	1 μ g/ml	14.8 \pm 3.9	17.2 \pm 1.7

5

Proliferation of two donor umbilical cord blood cultures further to induction with the CAHSP70 cloned fragments. Positive controls: C. albicans mannoproteic antigen (MP-F2) and Interleukin-2 (IL-2). Negative controls: no materials. Shown values represent average values \pm SD from 3 wells. For technical details, see table III legend and ref. 7.

10

Furthermore, immunoblotting experiments revealed the presence of anti-CAHSP70 antibodies in sera from adult wealthy humans, and in particular of the anti-CAHSP70/4 fragment (Fig. 8), suggesting that this fragment contains the immunodominant epitope. Taken together, lymphoproliferations human serum immunoblotting data suggest unequivocally that CAHSP70 is recognized by the immune system during the Candida usual colonization of healthy subjects.

15

20

Moreover, in immunoblotting on nitrocellulose filters, anti-CAHSP70 murine sera recognize more than one component of the HSP70 family from heat induced C. albicans extracted proteins (Fig. 5), thus showing that the expression product of caRLV130 insert is a C.

25

albicans protein which is expressed after the heat shock. According to the above results we named as CAHSP70 the C. albicans protein having the following properties: I) it comprises the aminoacid sequence coded by the caRLV130 insert; II) its gene maps on C. albicans chromosome 1 (having the highest molecular weight); III) its expression is induced by temperature shift; IV) it induces specific antibodies able to recognize cloned and purified fragments (subunits); V) it induces a lymphoproliferation in lymphomonocytic cultures from peripheral human blood. The relevant gene was named as cahsp70.

The CAHSP70 cloning, and its molecular and biochemical characterization, allows to develop a diagnostic molecular method based upon the amplification of DNA inserts corresponding to caRLV130, other than immunological studies of C. albicans 70 kDa heat shock protein expression. According to the caRLV130 insert sequence, we have synthesized oligonucleotides which were utilized for polymerase chain reaction (PCR) experiments, to analyze their ability to amplify DNA fragments which are homologous to C. albicans caRLV130 DNA. Two oligonucleotides (CA2-CA4) were chosen in the regions showing the minimal homology between the caRLV130 cDNA sequence and known HSP70 coding gene sequences (see Fig. 2 for the caRLV130 and YSCSSA1 sequence aligning, see Table V for the definition of minimal homology regions and Table VI for the sequence of oligonucleotides which were utilized for the assay).

The combination of CA2 (GAAATGAAAGATAAGATTGGTGCA) and CA3 (CCACAGTAAATTACCTATTTCTTCCTC) oligonucleotides is able to amplify DNA fragments having the expected size and a sequence specific of C. albicans DNA (Fig. 9A), whereas the assay sensitivity is shown in Fig. 9B by

using CA1 (ATGTCTAAAGCTGTTGGTATTG) and CA4
(CTGCACCAATCTTATCTTTCATTTACCATCATT) oligonucleotides.

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SEQUENCE LISTING

(1) GENERAL INFORMATION:

5 (i) APPLICANT:

(A) NAME: Istituto Superiore di Sanita'

(B) STREET: Viale Regina Elena 299

(C) CITY: Rome

(E) COUNTRY: Italy

10 (F) POSTAL CODE (ZIP): 00161

(A) NAME: Universita' degli Studi di Roma La Sapienza

(B) STREET: P.le Aldo Moro 5

(C) CITY: Rome

15 (E) COUNTRY: Italy

(F) POSTAL CODE (ZIP): 00184

(ii) TITLE OF INVENTION: Candida heath shock protein, gene and
uses thereof

20

(iii) NUMBER OF SEQUENCES: 2

(iv) COMPUTER READABLE FORM:

(A) MEDIUM TYPE: Floppy disk

25 (B) COMPUTER: IBM PC compatible

(C) OPERATING SYSTEM: PC-DOS/MS-DOS

(D) SOFTWARE: PatentIn Release #1.0, Version #1.30 (EPO)

(2) INFORMATION FOR SEQ ID NO: 1:

30

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 2001 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: single

35 (D) TOPOLOGY: linear

(ix) FEATURE:

(A) NAME/KEY: CDS

(B) LOCATION:1..1968

40

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 1:

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	Ala His Phe Ala Asn Asp Arg Val Glu Ile Ile Ala Asn Asp Gln Gly	
	20 25 30	
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	35 40 45	
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20	Val Phe Asp Ala Lys Arg Leu Ile Gly Arg Lys Phe Asp Asp Pro Glu	
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	Val Ile Asn Asp Ala Lys His Phe Pro Phe Lys Val Ile Asp Lys Ala	
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	145 150 155 160	
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	Leu Ser Ser Ser Ala Gln Thr Ser Ile Glu Ile Asp Ser Leu Tyr Glu	
	275 280 285	
35	GGT ATT GAC TTC TAC ACT TCA ATC ACC AGA GCC AGA TTT GAA GAA TTG	912
	Gly Ile Asp Phe Tyr Thr Ser Ile Thr Arg Ala Arg Phe Glu Glu Leu	
	290 295 300	
40	TGT GCT GAC TTG TTT AGA TCC ACT TTA GAT CCA GTT GGT AAA GTT TTA	960
	Cys Ala Asp Leu Phe Arg Ser Thr Leu Asp Pro Val Gly Lys Val Leu	
	305 310 315 320	
45	GCT GAT GCC AAG ATT GAT AAA TCT CAA GTT GAA GAA ATT GTC TTG GTT	1008
	Ala Asp Ala Lys Ile Asp Lys Ser Gln Val Glu Glu Ile Val Leu Val	
	325 330 335	
50	GGT GGG TCC ACC AGA ATT CCA AAG ATT CAA AAA TTG GTT TCT GAT TTC	1056
	Gly Gly Ser Thr Arg Ile Pro Lys Ile Gln Lys Leu Val Ser Asp Phe	
	340 345 350	

	TTT AAT GGT AAA GAA TTG AAT AAA TCT ATC AAC CCT GAT GAA GCT GTT	1104
	Phe Asn Gly Lys Glu Leu Asn Lys Ser Ile Asn Pro Asp Glu Ala Val	
	355 360 365	
5	GCT TAT GGT GCT GCT GTT CAA GCT GCC ATT TTA ACT GGT GAT ACT TCT	1152
	Ala Tyr Gly Ala Ala Val Gln Ala Ala Ile Leu Thr Gly Asp Thr Ser	
	370 375 380	
10	TCC AAG ACT CAA GAT ATT TTG TTA TTG GAT GTT GCT CCA TTG TCA TTA	1200
	Ser Lys Thr Gln Asp Ile Leu Leu Leu Asp Val Ala Pro Leu Ser Leu	
	385 390 395 400	
15	GGT ATT GAA ACT GCT GGT GGT ATC ATG ACC AAA TTG ATT CCA AGA AAT	1248
	Gly Ile Glu Thr Ala Gly Gly Ile Met Thr Lys Leu Ile Pro Arg Asn	
	405 410 415	
20	TCT ACT ATT CCA ACT AAG AAA TCA GAA ACT TTC TCC ACT TAT GCC GAT	1296
	Ser Thr Ile Pro Thr Lys Lys Ser Glu Thr Phe Ser Thr Tyr Ala Asp	
	420 425 430	
25	AAC CAA CCA GGT GTT TTG ATT CAA GTG TTT GAA GGT GAA AGA GCT AAA	1344
	Asn Gln Pro Gly Val Leu Ile Gln Val Phe Glu Gly Glu Arg Ala Lys	
	435 440 445	
30	ACT AAA GAT AAC AAC TTG TTG GGT AAA TTT GAA TTA TCT GGT ATT CCA	1392
	Thr Lys Asp Asn Asn Leu Leu Gly Lys Phe Glu Leu Ser Gly Ile Pro	
	450 455 460	
35	CCA GCT CCA AGA GGC GTC CCT CAA ATT GAA GTT ACT TTC GAT ATT GAT	1440
	Pro Ala Pro Arg Gly Val Pro Gln Ile Glu Val Thr Phe Asp Ile Asp	
	465 470 475 480	
40	GCT AAT GGT ATC TTG AAT GTT TCT GCT TTA GAA AAA GGT ACT GGT AAA	1488
	Ala Asn Gly Ile Leu Asn Val Ser Ala Leu Glu Lys Gly Thr Gly Lys	
	485 490 495	
45	ACT CAA AAG ATT ACT ATC ACC AAC GAT AAA GGT AGA TTA TCC AAA GAA	1536
	Thr Gln Lys Ile Thr Ile Thr Asn Asp Lys Gly Arg Leu Ser Lys Glu	
	500 505 510	
50	GAA ATT GAT AAA ATG GTT AGT GAA GCT GAA AAA TTC AAA GAA GAA GAT	1584
	Glu Ile Asp Lys Met Val Ser Glu Ala Glu Lys Phe Lys Glu Glu Asp	
	515 520 525	

	GAA AAG GAA GCT GCT AGA GTC CAA GCC AAG AAT CAA TTG GAA TCT TAT	1632
	Glu Lys Glu Ala Ala Arg Val Gln Ala Lys Asn Gln Leu Glu Ser Tyr	
	530 535 540	
5	GCT TAT TCA TTG AAA AAC ACA ATC AAT GAT GGT GAA ATG AAA GAT AAG	1680
	Ala Tyr Ser Leu Lys Asn Thr Ile Asn Asp Gly Glu Met Lys Asp Lys	
	545 550 555 560	
10	ATT GGT GCA GAT GAT AAA GAA AAA TTA ACT AAA GCC ATT GAT GAA ACT	1728
	Ile Gly Ala Asp Asp Lys Glu Lys Leu Thr Lys Ala Ile Asp Glu Thr	
	565 570 575	
15	ATT TCT TGG TTA GAT GCA TCT CAA GCT GCT TCT ACT GAA GAA TAC GAA	1776
	Ile Ser Trp Leu Asp Ala Ser Gln Ala Ala Ser Thr Glu Glu Tyr Glu	
	580 585 590	
20	GAT AAA CGT AAA GAA TTA GAA TCA GTT GCT AAT CCA ATC ATT AGT GGT	1824
	Asp Lys Arg Lys Glu Leu Glu Ser Val Ala Asn Pro Ile Ile Ser Gly	
	595 600 605	
25	GCT TAT GGT GCT GCC GGT GGC GCT CCA GGT GGT GCA GGC GGA TTC CCA	1872
	Ala Tyr Gly Ala Ala Gly Gly Ala Pro Gly Gly Ala Gly Gly Phe Pro	
	610 615 620	
30	GGT GCT ACT GGT GGT GAA TCT AGT GGA CCA ACT GTT GAA GAA GTT GAT	1920
	Gly Ala Gly Gly Phe Pro Gly Gly Ala Pro Gly Ala Gly Gly Pro Gly	
	625 630 635 640	
35	GGT GCT ACT GGT GGT GAA TCT AGT GGA CCA ACT GTT GAA GAA GTT GAT	1968
	Gly Ala Thr Gly Gly Glu Ser Ser Gly Pro Thr Val Glu Glu Val Asp	
	645 650 655	
	TAAATGAGGAAGAAATAGGTAATTTACTGTGG	2000

(2) INFORMATION FOR SEQ ID NO: 2:

(i) SEQUENCE CHARACTERISTICS:

- 40 (A) LENGTH: 656 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 2:

Met Ser Lys Ala Val Gly Ile Asp Leu Gly Thr Thr Tyr Ser Cys Val
 1 5 10 15
 5 Ala His Phe Ala Asn Asp Arg Val Glu Ile Ile Ala Asn Asp Gln Gly
 20 25 30
 Asn Arg Thr Thr Pro Ser Phe Val Ala Phe Thr Asp Thr Glu Arg Leu
 35 40 45
 Ile Gly Asp Ala Ala Lys Asn Gln Ala Ala Met Asn Pro Ala Asn Thr
 10 50 55 60
 Val Phe Asp Ala Lys Arg Leu Ile Gly Arg Lys Phe Asp Asp Pro Glu
 65 70 75 80
 Val Ile Asn Asp Ala Lys His Phe Pro Phe Lys Val Ile Asp Lys Ala
 85 90 95
 15 Gly Lys Pro Val Ile Gln Val Glu Tyr Lys Gly Glu Thr Lys Thr Phe
 100 105 110
 Ser Pro Glu Glu Ile Ser Ser Met Val Leu Thr Lys Met Lys Glu Ile
 115 120 125
 Ala Glu Gly Tyr Leu Gly Ser Thr Val Lys Asp Ala Val Val Thr Val
 20 130 135 140
 Pro Ala Tyr Phe Asn Asp Ser Gln Arg Gln Ala Thr Lys Asp Ala Gly
 145 150 155 160
 Thr Ile Ala Gly Leu Asn Val Leu Arg Ile Ile Asn Glu Pro Thr Ala
 165 170 175
 25 Ala Ala Ile Ala Tyr Gly Leu Asp Lys Lys Gly Ser Arg Gly Glu His
 180 185 190
 Asn Val Leu Ile Phe Asp Leu Gly Gly Gly Thr Phe Asp Val Ser Leu
 195 200 205
 Leu Ala Ile Asp Glu Gly Ile Phe Glu Val Lys Ala Thr Ala Gly Asp
 30 210 215 220
 Thr His Leu Gly Gly Glu Asp Phe Asp Asn Arg Leu Val Asn Phe Phe
 225 230 235 240
 Ile Gln Glu Phe Lys Arg Lys Asn Lys Lys Asp Ile Ser Thr Asn Gln
 245 250 255
 35 Arg Ala Leu Arg Arg Leu Arg Thr Ala Cys Glu Arg Ala Lys Arg Thr
 260 265 270
 Leu Ser Ser Ser Ala Gln Thr Ser Ile Glu Ile Asp Ser Leu Tyr Glu
 275 280 285
 Gly Ile Asp Phe Tyr Thr Ser Ile Thr Arg Ala Arg Phe Glu Glu Leu
 40

	290		295		300
	Cys Ala Asp Leu Phe Arg Ser Thr Leu Asp Pro Val Gly Lys Val Leu				
	305		310		315
	Ala Asp Ala Lys Ile Asp Lys Ser Gln Val Glu Glu Ile Val Leu Val				
5		325		330	335
	Gly Gly Ser Thr Arg Ile Pro Lys Ile Gln Lys Leu Val Ser Asp Phe				
		340		345	350
	Phe Asn Gly Lys Glu Leu Asn Lys Ser Ile Asn Pro Asp Glu Ala Val				
		355		360	365
10	Ala Tyr Gly Ala Ala Val Gln Ala Ala Ile Leu Thr Gly Asp Thr Ser				
		370		375	380
	Ser Lys Thr Gln Asp Ile Leu Leu Leu Asp Val Ala Pro Leu Ser Leu				
		385		390	395
	Gly Ile Glu Thr Ala Gly Gly Ile Met Thr Lys Leu Ile Pro Arg Asn				
15		405		410	415
	Ser Thr Ile Pro Thr Lys Lys Ser Glu Thr Phe Ser Thr Tyr Ala Asp				
		420		425	430
	Asn Gln Pro Gly Val Leu Ile Gln Val Phe Glu Gly Glu Arg Ala Lys				
		435		440	445
20	Thr Lys Asp Asn Asn Leu Leu Gly Lys Phe Glu Leu Ser Gly Ile Pro				
		450		455	460
	Pro Ala Pro Arg Gly Val Pro Gln Ile Glu Val Thr Phe Asp Ile Asp				
		465		470	475
	Ala Asn Gly Ile Leu Asn Val Ser Ala Leu Glu Lys Gly Thr Gly Lys				
25		485		490	495
	Thr Gln Lys Ile Thr Ile Thr Asn Asp Lys Gly Arg Leu Ser Lys Glu				
		500		505	510
	Glu Ile Asp Lys Met Val Ser Glu Ala Glu Lys Phe Lys Glu Glu Asp				
		515		520	525
30	Glu Lys Glu Ala Ala Arg Val Gln Ala Lys Asn Gln Leu Glu Ser Tyr				
		530		535	540
	Ala Tyr Ser Leu Lys Asn Thr Ile Asn Asp Gly Glu Met Lys Asp Lys				
		545		550	555
	Ile Gly Ala Asp Asp Lys Glu Lys Leu Thr Lys Ala Ile Asp Glu Thr				
35		565		570	575
	Ile Ser Trp Leu Asp Ala Ser Gln Ala Ala Ser Thr Glu Glu Tyr Glu				
		580		585	590
	Asp Lys Arg Lys Glu Leu Glu Ser Val Ala Asn Pro Ile Ile Ser Gly				
		595		600	605
40	Ala Tyr Gly Ala Ala Gly Gly Ala Pro Gly Gly Ala Gly Gly Phe Pro				

	610					615						620					
	Gly	Ala	Gly	Gly	Phe	Pro	Gly	Gly	Ala	Pro	Gly	Ala	Gly	Gly	Pro	Gly	
	625					630					635					640	
	Gly	Ala	Thr	Gly	Gly	Glu	Ser	Ser	Gly	Pro	Thr	Val	Glu	Glu	Val	Asp	
5					645					650						655	

Claims

1. A nucleic acid comprising a nucleotide sequence coding the protein having the amino acid sequence of SEQ
5 ID No.2 or parts thereof.

2. A nucleic acid comprising a nucleotide sequence with at least a 65% homology with the nucleotide sequence of SEQ ID No.1 or parts thereof.

3. A nucleic acid according to claim 2 comprising
10 the nucleotide sequence of SEQ ID No.1 or parts thereof.

4. Composition comprising a nucleic acid according to any of claims 1 to 3.

5. Use of the nucleic acid according to any of claims 1 to 3 for oligonucleotide probes to be used in
15 diagnosis and typing of Candida and Candida related pathologies.

6. Oligonucleotide having a sequence comprised in SEQ ID No. 1 to be used for PCR (polymerase chain reaction) to detect the presence in biological and/or environment
20 samples either of C. albicans or of other Candida species or of yeast-like related microorganisms comprising said gene and/or in a labeled form (radionuclides, biotin, enzymes, etc.) to detect the presence in biological and/or environment samples either of C. albicans or of
25 other related and/or for the C. albicans or related species typing and/or diagnosis and/or as potential antibiotic and/or chemiotherapeutic targets, or antisense RNA active for Candida species and/or yeast-like related microorganisms coding an homologous sequence.

30 7. Polypeptide having the aminoacid sequence comprised in the SEQ ID No.1, or having at least a 50% homology with SEQ ID No. 1 or fragments, and/or functional and immunologic homologous thereof.

```

atg tct aaa gct gtt ggt att gat tta ggt aca acc tat tct tgt gtt gct cat ttt gcc aat gat aga gtt gaa att att gct aat gat caa ggt aat aga act acc cct
M S K A V G I D L G T T Y S C V A H F A N D R V E I I A N D Q G N R T T P>
100

tca ttt gtt gcc ttc act gat act gaa aga ttg att ggt gat gct gcc aag aat caa gct gct atg aac cca gca aac act gtt ttc gat gct aaa cgt tta att ggg aga
S F V A F T D T E R L I G D A A K N Q A A M N P A N T V F D A K R L I G R>
200

aaa ttt gat gat cca gaa gtt ata aat gat gct aaa cat ttc cca ttt aaa gtc att gat aaa gca ggt aaa cca gtg att caa gtt gaa tat aaa ggt gaa act aaa act
K F D D P E V I N D A K H F P F K V I D K A G K A P V I Q V E Y K G E T K T>
300

ttt tca cca gaa gaa att tct tca atg gtt tta aca aaa atg aaa gaa att gct gaa ggt tat ttg ggt tct act gtt aaa gat gct gtt gtt act gtt cca gct tat ttc
F S P E E I S S M V L T K M K E I A E G Y L G S T V K D A V V T V P A Y F>
400

aat gat tct tca aga cca gcc acc aaa gat gct ggt act att gct ggt ttg aat gtt tta aga att aat aat gaa cct act gct gcc att gct tat ggt tta gat aaa
N D S Q R Q A T K D A G T I A G L N V L R I I N E P T A A I A Y G L D K>
500

aaa ggt tcc aga ggt gaa cat aat gtt tta att ttc gat ttg ggt ggt ggt act ttt gat gtt tca tta tta gcc att gat gaa ggt att ttc gaa gtt aaa gcc act gct
K G S R G E H N V L I F D L G G G T F D V S L L A I D E G I F E V K A T A>
600

ggt gat act cat ttg ggt ggt gaa gat ttt gat aac aga tta gtc aac ttc ttt att caa gaa ttc aag aga aag aac aag aaa gat att tcc acc aac caa aga gct tta
G D T H L G G E D F D N R L V N F F I Q E F K R K N K K D I S T N Q R A L>
700

aga aga tta aga act gct tgt gaa aga gcc aag aga act ttg tct tct gct caa acc tca att gaa att gat tcc tta tat gaa ggt att gac ttc act tca atc
R R I, R T A C E R A K R T L S S S A Q T S I E I D S L Y E G I D F Y T S I>
800

acc aga gcc aga ttt gaa gaa ttg gct gac ttg ttt aga tcc act tta gat cca gtt ggt aaa gtt tta gct gat gcc aag att gat aaa tct caa gtt gaa gaa att
T R A R F E L C A D L F R S T L D P V G K V L A D A K I D K S Q V E E I>
900

```

FIG. 1

FIG. 1(cont)

FIG. 2

[illegible]

FIG. 2 (cont)

FIG. 2 (cont)

mskavgldlgttyscvahfaukrvel lairdgnrt tpsfvafctder l l gdaaknqaampant v f d a k r l l g r k d d i p e v l n d a k h f p f k v i d k a g k p v l q v e y k g e t k t f a p e e l s a m v l t k m k e l a e g y l g e t v k d a v v t v p a y f n d
10 20 30 40 50 60 70 80 90 100 110 120 130 140 150
MSKAVGIDLGHTYSCVAHFANDRVDI IANQQCNRTTPSFVAFDTIERLIGDAANKQMAANPSNVFEDAKRLIGRNFNQPEVQATMKIIFPEKLLIDVDCKPOIQVEFKGETKNFPEQI SSMVLGKHKETASVYLGAKVNDAAVTVPAVRND>
mskavgldlgttyscvahfaukrvel lairdgnrt tpsfvafctder l l gdaaknqaampant v f d a k r l l g r k d d i p e v l n d a k h f p f k v i d k a g k p v l q v e y k g e t k t f a p e e l s a m v l t k m k e l a e g y l g e t v k d a v v t v p a y f n d
200 300
sqratkdagtlaglnvrl l l n e p t a a a l a y g l d k k g s r g e h n v l l f d l g g g t f d v e l l a l d e g l f e v k a t a g d t h l g g e d f d n r l v n f f l g e f k r k u k k l a t n q r a l r r l r t a c e r a k r t l a s s a q t a l e l d a l y e g l d f y t s l t a r
160 170 180 190 200 210 220 230 240 250 260 270 280 290
SQRQATKDA GTT IAGLNVLRI INEPTAAAIAYGLDKKG-KEEH-VLI FDLGGCTFDVSLLFIEDGIFEVKATACDTHILGGEDEFDRILVNIIFIOEFKRKNKDLSTINQRAI RRLRTACERAKRTLSSSAQGSVEIDSLFEGIDFVTSITRAR>
sqratkdagtlaglnvrl l l n e p t a a a l a y g l d k k g s r g e h n v l l f d l g g g t f d v e l l a l d e g l f e v k a t a g d t h l g g e d f d n r l v n f f l g e f k r k u k k l a t n q r a l r r l r t a c e r a k r t l a s s a q t a l e l d a l y e g l d f y t s l t a r
400 500 600
feelcadl l f r e t l d p v g k v l a d a k l d k s q v e e l v l v g g s t r i p k l q k l v e d f f n g k e l n k s l n p d e a v a y g a a v q a a l l t g d t e s k t q d l l l d v a p l a l g l e t a g g l m t k l l p r n e t l p c k k e e t f s t y a d n q p g v l l q v f e g e r a k t k
300 310 320 330 340 350 360 370 380 390 400 410 420 430 440
FEELCADLFRSTLDPVEKVL RDAKLDK SQVDEIVLVGGS TRIPK VQKLVTDYFRNGKEFNRSINPDEAVAYGA VQAALITGDESSKTCOLL LLDVAPLSLGLETAGGVMTKLI PRNSTISTRKFEIFSTYADNQPGVL IQVFEGERAKTR>
feelcadl l f r e t l d p v g k v l a d a k l d k s q v e e l v l v g g s t r i p k l q k l v e d f f n g k e l n k s l n p d e a v a y g a a v q a a l l t g d t e s k t q d l l l d v a p l a l g l e t a g g l m t k l l p r n e t l p c k k e e t f s t y a d n q p g v l l q v f e g e r a k t k
500 600
dnmllgkfelsglppaprgvqjlevtfldldangl lnvealekgtgktqklt l tndkgr l s k e e l d k m v s e a e k f k e e d e k e a a r v q a k n q l e s y a y s l k n t l n d g e m k d k l g a d d k e k l t c k a l d e t l s w l d a s q a a s t e e y e d k r k e l e s
450 460 470 480 490 500 510 520 530 540 550 560 570 580 590
DNMLLGKFELSGIPPA TRGVQJLEVTFDVSNGI LNVS AVEKGTGKSHIKITITNDKGRLSKEDI EKMWAEKEKEEDESORIASKNQLES IAYS LKNRTISEAG--DKLEQADKDTVTYKAEETI SMLDSNTYASKEEFDOKLKEIQD>
dnmllgkfelsglppaprgvqjlevtfldldangl lnvealekgtgktqklt l tndkgr l s k e e l d k m v s e a e k f k e e d e k e a a r v q a k n q l e s y a y s l k n t l n d g e m k d k l g a d d k e k l t c k a l d e t l s w l d a s q a a s t e e y e d k r k e l e s
600 610 620 630 640
VANPILSGAYGAAGGAPGAGG fpgagg fpxjapagagrggagtgessgptveevd*
IANPIMSKLIY-OAGCATGGAAG-CATGCGFPGCAPPAPEAEGETVEE>
vanpilisgayaagagagagg fpgagg fpgapagagrggagtgagg

FIG. 3

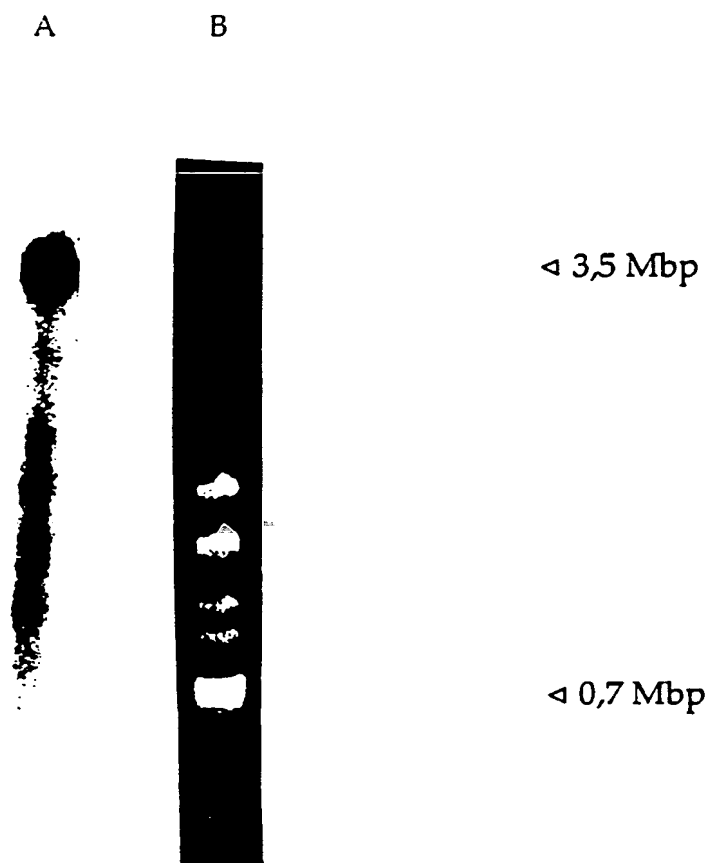


FIG. 4

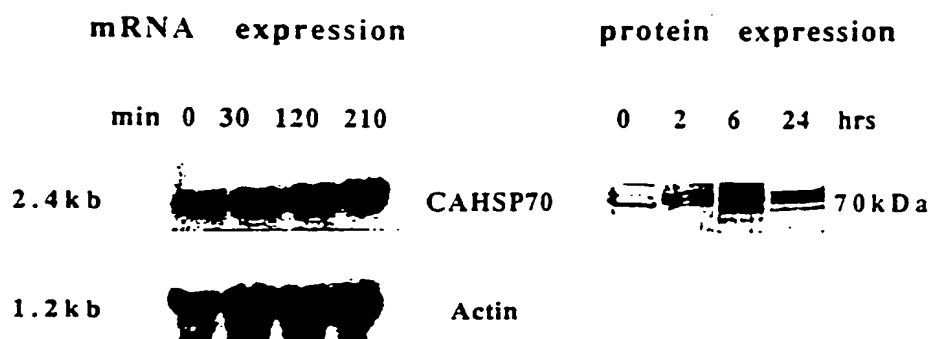
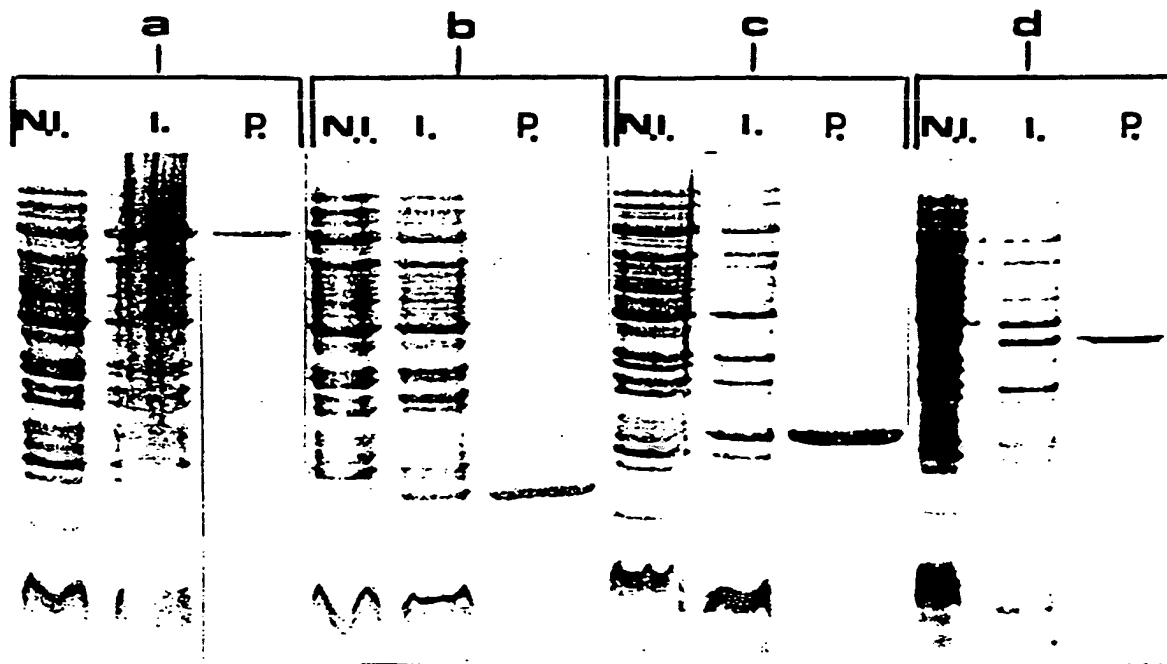
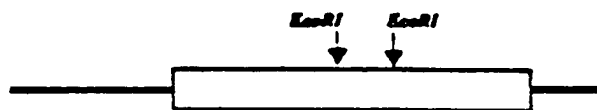


FIG. 5

A



B



CAHSP70	N-6xHis-	<div style="border: 1px solid black; padding: 2px; display: inline-block;">a</div>	-C	<71kDa
CAHSP70/2		N-6xHis-	<div style="border: 1px solid black; padding: 2px; display: inline-block;">b</div>	-C <21kDa
CAHSP70/3	N-6xHis-	<div style="border: 1px solid black; padding: 2px; display: inline-block;">c</div>	-C	<28kDa
CAHSP70/4	N-6xHis-	<div style="border: 1px solid black; padding: 2px; display: inline-block;">d</div>	-C	<39kDa

FIG. 6

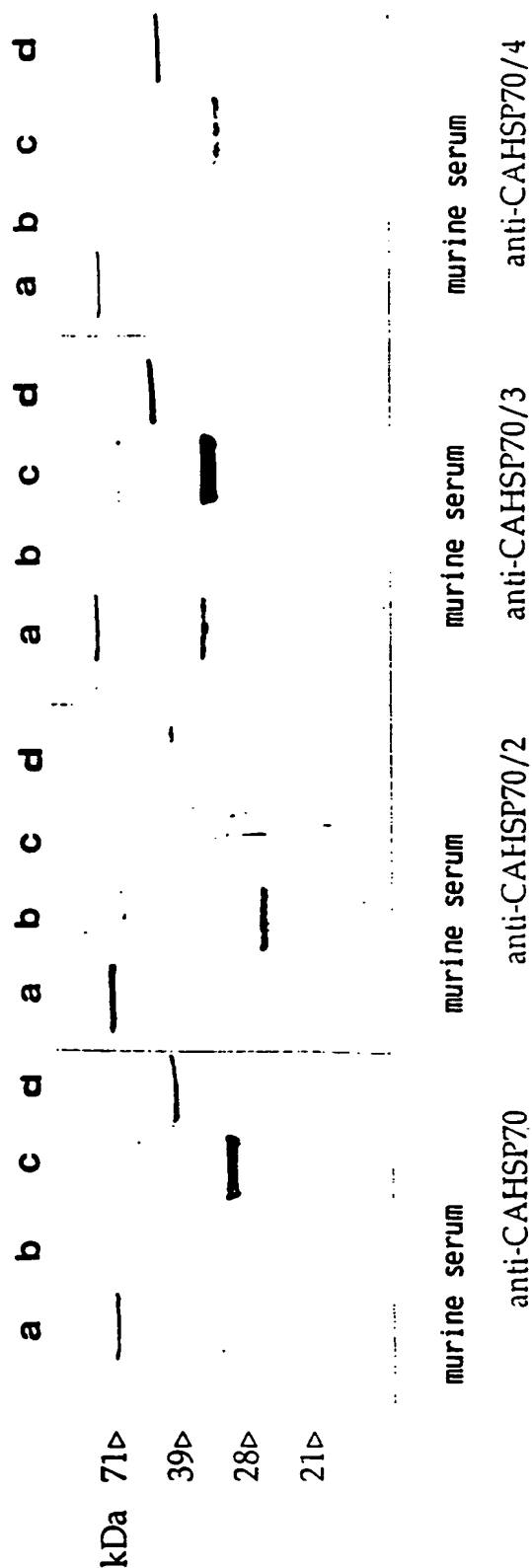


FIG. 7

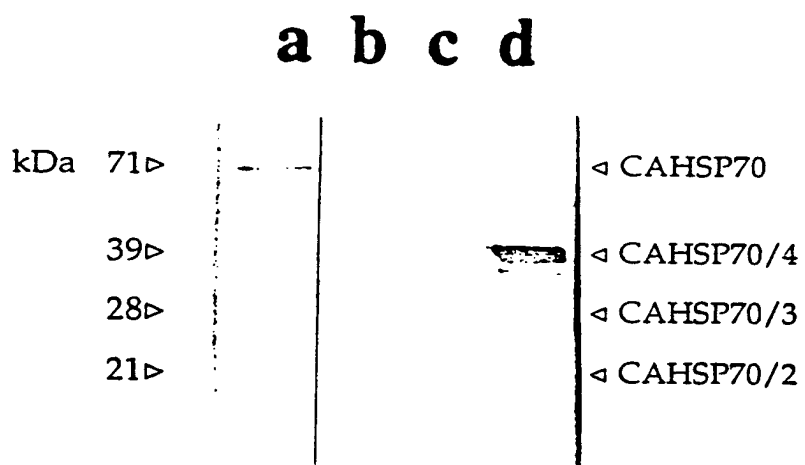


FIG. 8

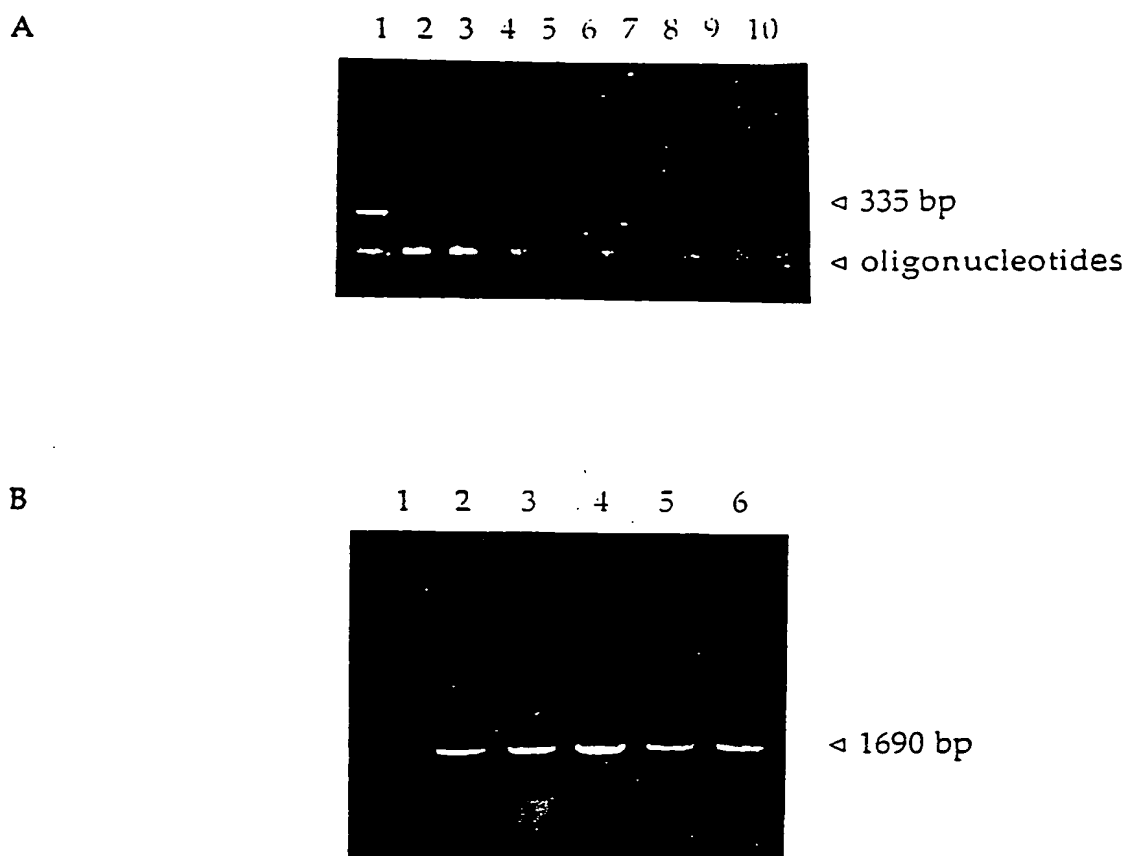


FIG. 9

INTERNATIONAL SEARCH REPORT

International Application No.

PCT 96/00097

A. CLASSIFICATION OF SUBJECT MATTER

IPC 6 C12N15/11 C12Q1/68 C07K14/40 C07K16/14 G01N33/569
A61K39/00

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 C12N C12Q C07K G01N A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	NUCLEIC ACIDS RESEARCH; vol. 17, no. 2, 25 January 1989, OXFORD GB, pages 805-806, XP002011023 M.R.SLATER AND E.A.CRAIG: "The SSA1 and SSA2 genes of the yeast <i>Saccharomyces cerevisiae</i> " see figure 1	1-4,7,8
A	--- EP,A,0 406 029 (J.P.BURNIE AND R.C.MATTHEWS) 2 January 1991 see the whole document --- -/-	9-12

☒ Further documents are listed in the continuation of box C.☒ Patent family members are listed in annex.

* Special categories of cited documents:

- * "A" document defining the general state of the art which is not considered to be of particular relevance
- * "E" earlier document but published on or after the international filing date
- * "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- * "O" document referring to an oral disclosure, use, exhibition or other means
- * "P" document published prior to the international filing date but later than the priority date claimed

* "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

* "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

* "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

* "A" document member of the same patent family

Date of the actual completion of the international search

19 August 1996

Date of mailing of the international search report

28.08.96

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax (+31-70) 340-3016

Authorized officer

Cupido, M

INTERNATIONAL SEARCH REPORT

International Application No.

PCT/IT 96/00097

C(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P,X	INFECTION AND IMMUNITY, vol. 63 , no. 10, October 1995, WASHINGTON US, pages 4039-4045, XP002011024 R.LA VALLE ET AL.: "Molecular cloning and expression of a 70-kilodalton heat shock protein of Candida albicans" see the whole document ---	1-12
A	JOURNAL OF MOLECULAR EVOLUTION, vol. 38, no. 1, January 1994, pages 1-17, XP000578395 W.R.BOORSTEIN ET AL.: "Molecular evolution of the HSP70 multigene family" see table 3 -----	1-4,7,8

FURTHER INFORMATION CONTINUED FROM PCT/ISA/210

Remark : Although claim 12, insofar as it relates to in vivo uses, is directed to a method of treatment of the human/animal body the search has been carried out and based on the alleged effects of the composition.

INTERNATIONAL SEARCH REPORT

International application No.

PCT/IT 96/00097

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:
Please see Further Information sheet enclosed.
2. ☐ Claims Nos.:
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No.

IT 96/00097

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
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